

Effectum Medical AG

assists in the development of innovative medical devices and technologies throughout the certification and life cycle management process. Acting as legal manufacturer and by offering our Plug & Play (e)QMS, we speed up your time to market and ensure compliance and excellence in every stage of development.



AI/ML DEVICE CERTIFICATION



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Al and ML Device Certification for Healthcare Navigate Al/ML Integration with Confidence

Incorporating Machine Learning (ML) and Artificial Intelligence (AI) into medical devices brings transformative potential, but it also requires navigating a complex regulatory environment. Effectum Medical provides expert guidance to ensure your devices meet all relevant compliance standards, including the European AI Act, MDR, and IVDR.





Our Comprehensive AI/ML Compliance Services

AI Readiness & Compliance Checks

- Gap Analyses & Al Readiness: We review your documentation to identify any gaps, ensuring it aligns with Al Act requirements.
- Dual or Single Regulation Assessments:
 Tailored checks to meet both AI Act and MDR/IVDR compliance, providing peace of mind at every stage.

CE Certification Support

Expert Guidance: Our team assists in achieving CE certification, offering specialized support for ML/AI-based software to help you navigate the approval process efficiently.

Technical Documentation Templates

Streamlined Preparation: Access customizable templates that simplify the preparation of technical files, helping you maintain compliance while saving time.

AI Act Compliance Solutions

 Harmonized Compliance Strategy: Our established processes ensure that your systems meet both AI Act and MDR/IVDR requirements efficiently and effectively.





From development through to post-market monitoring, we support your ML/AI device across its entire lifecycle:

- Pre-market Conformity Assessments: Determine if a notified body conformity assessment is necessary or if internal controls will apply. Our team will also assist in preparing technical documentation in line with both AI Act and MDR/IVDR standards.
- **Clinical Evaluation Support:** We help align clinical evaluations with ML/Al-specific regulatory requirements.
- **Continuous Monitoring Solutions:** Ensure your device remains compliant as it evolves in real-world use, meeting ongoing monitoring and regulatory requirements.

Preparing for the Future of Healthcare AI Compliance

Developing a robust ML/AI device can take 1-2 years, with conformity assessments adding another year. To meet the August 2027 deadline for high-risk applications under the EU AI Act, it's critical to start early.



Why Choose Effectum Medical?

- **Dual Expertise:** Specialists in AI regulation and medical device compliance.
- Efficiency: Streamlined processes that align with AI Act and MDR/IVDR.
- **Future-Proofing:** Preparedness for upcoming regulatory changes, ensuring your product's compliance and adaptability.

Don't let regulatory requirements stall your healthcare AI innovations. **Partner with Effectum Medical** to turn compliance into a growth opportunity and build patient trust.

Contact us for a personalized consultation to prepare your ML/AI models for the European market.







Outsourced Legal Manufacturer:

Streamline market access and compliance with our trusted legal manufacturer services. Need speed? Ask about our fast-track option!



Plug & Play (e)QMS:

Accelerate your launch with a ready-made EN ISO 13485, MDR/IVDR, and FDA-compliant quality management system.



Plug-and-Play eQMS Bootcamp:

Flexible training to empower self-management of your QMS.



Rent an Expert:

Add quality management and regulatory expertise to your team as needed.



Regulatory Opinions:

Access expert insights for global compliance and market strategy.



AI/ML Device Certification:

Certification services for AI and machine learning-based medical devices.

For more information visit: www.effectummedical.com



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